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                 CA/CAplus kind code changes for Chinese patents
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                 INPADOCDB/INPAFAMDB: Enhancements to the US national
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                 patent classification.
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      7 NOV 03
                 New format for Korean patent application numbers in
                 CA/CAplus increases consistency, saves time.
         NOV 04
                 Selected STN databases scheduled for removal on
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                 December 31, 2010
         NOV 18
                 PROUSDDR and SYNTHLINE Scheduled for Removal
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      9
                 December 31, 2010 by Request of Prous Science
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                 Higher System Limits Increase the Power of STN
NEWS 10
                 Substance-Based Searching
NEWS 11
         NOV 24
                 Search an additional 46,850 records with MEDLINE
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         DEC 14 New PNK Field Allows More Precise Crossover among STN
                 Patent Databases
         DEC 18 ReaxysFile available on STN
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         DEC 21
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         DEC 22 Value-Added Indexing Improves Access to World Traditional
                 Medicine Patents in CAplus
                 The new and enhanced DPCI file on STN has been released
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         JAN 24
NEWS 17 JAN 26
                 Improved Timeliness of CAS Indexing Adds Value to
                 USPATFULL and USPAT2 Chemistry Patents
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         JAN 26
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                 other enhancements improve searching in STN reload of
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NEWS 19
         JAN 28
                 CABA will be updated weekly
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         FEB 23
                 PCTFULL file on STN completely reloaded
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         FEB 23
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                 Qualified Customers
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         FEB 25
                 LPCI will be replaced by LDPCI
NEWS 23
         MAR 07
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                 Numbers in the USPAT and IFI Database Families is Now
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COST IN U.S. DOLLARS

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56 FILES IN THE FILE LIST IN STNINDEX

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  - 0\* FILE ANTE
  - 0\* FILE AQUALINE
  - 0\* FILE BIOENG
  - 0\* FILE BIOTECHABS
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  - 1 FILE CAPLUS
  - 0\* FILE CEABA-VTB
  - 0\* FILE CIN
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    - 0\* FILE FROSTI 0\* FILE FSTA
    - 1 FILE IFIPAT
    - 0\* FILE KOSMET
  - 38 FILES SEARCHED...
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    - 0\* FILE PASCAL
    - 3 FILE USPATFULL
    - 0\* FILE WATER
  - 3 FILES HAVE ONE OR MORE ANSWERS, 56 FILES SEARCHED IN STNINDEX
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- => file caplus ifipat uspatfull

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FILE 'USPATFULL' ENTERED AT 03:02:14 ON 28 MAR 2011 CA INDEXING COPYRIGHT (C) 2011 AMERICAN CHEMICAL SOCIETY (ACS)

=> s 11 L2 5 L1

=> dup rem 12
PROCESSING COMPLETED FOR L2

L3 4 DUP REM L2 (1 DUPLICATE REMOVED)

=> d 13 1-4

L3 ANSWER 1 OF 4 IFIPAT COPYRIGHT 2011 IFI on STN DUPLICATE 1

AN 11892997 IFIPAT; IFIUDB; IFICDB

TI Use of Probiotic Bacteria in the Treatment of Infection; use of a live culture of a non-pathogenic food-grade probiotic bacterium (lactococcus or lactobacillus strains of lactic acid bacteria) in the treatment of infectious diseases, may be a localised infection of the skin, including an infected wound, a urinary tract infection or mastitis

IN Hallahan Stephen (IE); Ross Paul (IE)

PA Unassigned Or Assigned To Individual (68000)

PPA Teagasc The Agriculture and Food Development Authority IE PPA University College Cork National University Or (Probable)

PI US 20080233091 Ā1 20080925

AI US 2004-576010 20041015 (10)

WO 2004-IE143 20041015

20070806 PCT 371 date 20070806 PCT 102(e) date

PRAI IE 2003-773 20031017 FI US 20080233091 20080925

DT Utility; Patent Application - First Publication

FS CHEMICAL

APPLICATION

ED Entered STN: 26 Sep 2008 Last Updated on STN: Jan 2011

CLMN 24

PΑ

L3 ANSWER 2 OF 4 USPATFULL on STN

AN 2008:200899 USPATFULL

TI Enterococcus Antigens

IN Meinke, Andreas, Pressbaum, AUSTRIA Nagy, Eszter, Vienna, AUSTRIA Hanner, Markus, Vienna, AUSTRIA

Gelbmann, Dieter, Andau, AUSTRIA
Intercell AG, Vienna, AUSTRIA (non-U.S. corporation)

PI US 20080175856 A1 20080724 AI US 2004-558119 A1 20040526 (10)

WO 2004-EP5664 20040526

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20051123 PCT 371 date
       EP 2003-450137
PRAI
                                20030530
       Utility
DТ
FS
       APPLICATION
LN.CNT 5615
INCL
       INCLM: 424/190.100
NCL
       NCLM:
             424/190.100
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       IPCI
              A61K0039-02 [I,A]; A61P0037-00 [I,A]
       IPCR
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              G01N0033-569 [I,A]
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     ANSWER 3 OF 4 USPATFULL on STN
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       2007:113675 USPATFULL
ΑN
ΤI
       Antagonistic properties of reef fish microflora
       Bruno, Cynthia K., Washington, DC, UNITED STATES
ΙN
РΤ
       US 20070098745
                         A1 20070503
                           A1
ΑI
       US 2006-589301
                                20061030 (11)
       Continuation-in-part of Ser. No. WO 2005-US15063, filed on 2 May 2005,
RLI
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       US 2004-566600P
                               20040430 (60)
PRAI
       Utility
DT
       APPLICATION
FS
LN.CNT 2252
INCL
       INCLM: 424/234.100
       INCLS: 424/243.100; 435/252.100
       NCLM: 424/234.100
NCL
       NCLS: 424/243.100; 435/252.100
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     2005:341762 CAPLUS
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     Use of Probiotic Bacteria in the Treatment of
ΤI
ΙN
     Ross, Paul; Hallahan, Stephen
     Teagasc the Agriculture and Food Development Authority, Ire.; University
PA
     College Cork
SO
     PCT Int. Appl.
     CODEN: PIXXD2
DТ
     Patent
     English
LA
FAN.CNT 1
     PATENT NO.
                        KIND
                                 DATE APPLICATION NO. DATE
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             GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI,
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IE 2003000773 A1
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20060705 EP 2004-770417
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                         A1
                                                                     20041015
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                         B1
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     AT 439851
                         Т
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                         A1
                               20080925
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PRAI IE 2003-773
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     WO 2004-IE143
                          W
                                20041015
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              THERE ARE 2 CAPLUS RECORDS THAT CITE THIS RECORD (2 CITINGS)
RE.CNT 10
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              ALL CITATIONS AVAILABLE IN THE RE FORMAT
=> d 13 2
     ANSWER 2 OF 4 USPATFULL on STN
1.3
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ΑN
ТΤ
       Enterococcus Antigens
ΙN
       Meinke, Andreas, Pressbaum, AUSTRIA
       Nagy, Eszter, Vienna, AUSTRIA
       Hanner, Markus, Vienna, AUSTRIA
       Gelbmann, Dieter, Andau, AUSTRIA
       Intercell AG, Vienna, AUSTRIA (non-U.S. corporation)
PΑ
       US 20080175856 A1 20080724 US 2004-558119 A1 20040526 (10)
PΙ
ΑI
       WO 2004-EP5664
                               20040526
                                20051123 PCT 371 date
PRAI
      EP 2003-450137
                               20030530
DT
       Utility
FS
      APPLICATION
LN.CNT 5615
INCL
       INCLM: 424/190.100
NCL
       NCLM: 424/190.100
IPC
       IPCI
              A61K0039-02 [I,A]; A61P0037-00 [I,A]
              A61K0039-02 [I,C]; A61K0039-02 [I,A]; A61K0039-00 [N,C*];
              A61K0039-00 [N,A]; A61P0037-00 [I,C]; A61P0037-00 [I,A];
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              G01N0033-569 [I,A]
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
=> d 13 2 kwic
     ANSWER 2 OF 4 USPATFULL on STN
       Enterococci are gram-positive bacteria that are normal
SUMM
       inhabitants of the alimentary tract of humans and animals. They have
       been recognized as a cause of. . .
SUMM
       . . E. faecium is unique because it is commonly used in production
       of fermented foods, and is also used as a probiotic bacterium. In recent years, E. faecium has been less acceptable as a food
       fermentation organism because of concern that this bacterium may be an
       intermediate host for spreading of antibiotic resistance to
       bacteria involved in human infections. Despite these concerns,
       E. faecium is still amongst the most common bacteria found in
       foods fermented by lactic acid bacteria. Many isolates of E.
       faecium have been shown to produce bacteriocins (antimicrobial peptides)
       that are able to kill or inhibit. . . been chosen as starter cultures
```

in the production of fermented food. Recently, enterocins have been implemented successfully in treatment of mastitis in cattle. Infection caused by the genus Enterococcus include a) bacteremia, b) SUMM urinary tract infections c) endophthalmitis, d) endocarditis and also wound and intra-abdominal infections. Approximately 3/4 of the infections are caused by. SUMM b) Urinary Tract Infections SUMM Enterococci have been estimated to account for 110,000 urinary tract infections (UTI) annually in the United States. A few studies have been aimed at understanding the interaction of enterococci. SUMM . . . select for appropriate screening reagents, a series of immunoassays (mainly ELISA and immunoblotting) were performed with bacterial lysate and culture supernatant proteins to measure anti-E. faecalis IgG antibody levels. Sera from high titer individuals were included in the genomic-based antigen identification. SUMM . . most important diseases, which can be inflicted by the two pathogens is presented below. S. aureus causes mainly nosocomial, opportunistic infections: impetigo, folliculitis, abscesses, boils, infected lacerations, endocarditis, meningitis, septic arthritis, pneumonia, osteomyelitis, scalded skin syndrome (SSS), toxic shock syndrome. E. faecalis causes mainly infections which are not highly toxigenic, highly invasive, or highly infectious by most measures. They do, nevertheless, cause a substantial amount of human disease such as bacteremia, urinary tract infections, endocarditis and intra-abdominal infections. . . a serum collection, which has been tested against antigenic SUMM compounds of E. faecalis, such as whole cell extracts and culture supernatant proteins. Preferably, 2 distinct serum collections are used: 1. With very stable antibody repertoire: normal adults, clinically healthy people, who. . induced acutely by the presence of the pathogenic organism: patients with acute disease with different manifestations (e.g. E. faecalis endocarditis, urinary tract infection and bacteraemia). Sera have to react with multiple enterococci-specific antigens in order to be considered hyperimmune and therefore. . . . against secreted proteins are beneficial in neutralisation of SUMM their function as toxin or virulence component. It is also known that bacteria communicate with each other through secreted proteins. Neutralizing antibodies against these proteins will interrupt growth-promoting cross-talk between or within enterococcal. SUMM . . . mechanisms. Inducing high affinity antibodies of the opsonic and neutralizing type by vaccination helps the innate immune system to eliminate bacteria and toxins. This makes the method according to the present invention an optimal tool for the identification of enterococcal antigenic. . . SUMM The skin and mucous membranes are formidable barriers against invasion by enterococci. However, once the skin or the mucous membranes are breached the first line of non-adaptive cellular defence begins its co-ordinate action through complement and phagocytes, especially the polymorphonuclear leukocytes (PMNs). These cells can be regarded as the cornerstones in eliminating invading bacteria. As enterococci are primarily extracellular pathogens, the major anti-enterococcal adaptive response comes from the humoral arm of the immune system,. . . to activated C3b. After opsonization, enterococci are phagocytosed and killed. Antibodies bound to specific antigens on the cell surface of bacteria serve as ligands for the attachment to PMNs and to promote phagocytosis. The very same antibodies bound to the adhesins. . . antigens as provided by the present invention is thus well suited to identify those that will lead to protection against infection in an animal model or in humans. SUMM . . of the invention can be synthetically produced by conventional

peptide synthesizers. Mature proteins can be expressed in mammalian cells, yeast, bacteria, or other cells under the control of appropriate promoters. Cell-free translation systems can also be, employed to produce such proteins. . .

SUMM . . a disease, most preferably for the diagnosis of a diseases related or linked to the presence or abundance of Gram-positive bacteria, especially bacteria selected from the group comprising enterococci, staphylococci and lactococci. More preferably, the microorganisms are selected from the group comprising Streptococcus.

SUMM . may be used as an antigen for vaccination of a host to produce specific antibodies which protect against invasion of bacteria , for example by blocking adherence of bacteria to damaged tissue. Examples of tissue damage include wounds in skin or connective tissue caused e.g. by mechanical, chemical or. .

SUMM . . described above, the compositions of this invention may be used generally as a wound treatment agent to prevent adhesion of bacteria to matrix proteins exposed in wound tissue and for prophylactic use in dental treatment as an alternative to, or in.

DETD Total bacterial lysate: Bacteria were grown overnight in BHI (Brain-heart Infusion) and lysed by repeated freeze-thaw cycles: incubation on dry ice/ethanol-mixture until frozen (1 min), then thawed at  $37^{\circ}$  C. (5 min): repeated 3 times. This was followed by sonication and collection of supernatant by centrifugation (3,500 rpm, 15 min, 4° C.).

DETD Culture supernatant: After removal of bacteria by centrifugation, the supernatant of overnight grown bacterial cultures was precipitated with ice-cold ethanol by mixing 1 part supernatant with 3 parts absolute ethanol and incubated overnight at -20° C. Precipitates were collected by centrifugation (2,600 g, for 15. . .

DETD Total bacterial lysate and culture supernatant samples were prepared from in vitro grown E. faecalis strain V583. 10 to 25  $\mu g$ total protein/lane was separated by. .

DETD . . . immune assays. Primary characterization was done by ELISA using two different antigen preparations, such as total bacterial extract and culture supernatant proteins prepared from E. faecalis V583 strain. Representative experiments are shown in FIG. 1 using sera from the healthy adult. . .

DETD Preparation of enterococcal genomic DNA. 50 ml Brain heart infusion (BHI) medium was inoculated with Enterococcus faecalis V583 bacteria from a frozen stab and grown with aeration and shaking for 18 h at 37° C. The culture was then harvested, centrifuged with  $1,600\times g$  for 15 min and the supernatant was removed. Bacterial pellets were washed 3× with PBS and carefully re-suspended in 0.5 ml of Lysozyme solution (100 mg/ml).. . solution was incubated for 20 min at 4  $^{\circ}$  C. The extract was pelleted in a microfuge (13,000 rpm) and the supernatant transferred into a new tube. The solution was extracted with PhOH/CHCl.sub.3/IAA (25:24:1) and with CHCl.sub.3/IAA (24:1). DNA was precipitated at. .

=> FIL STNGUIDE COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 30.47 33.54

FULL ESTIMATED COST

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0\* FILE KOSMET

0\* FILE NTIS

0\* FILE PASCAL

3 FILE USPATFULL

0\* FILE WATER

L1 QUE SKIN(P)INFECT? AND PROBIOTIC AND BACTERIA AND (SUPERNATANT

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4 DUP REM L2 (1 DUPLICATE REMOVED)

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L3

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=> index bioscience
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FULL ESTIMATED COST

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56 FILES IN THE FILE LIST IN STNINDEX

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  - 1 FILE BIOENG
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  - 1 FILE MEDLINE
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  - 2 FILE SCISEARCH
  - 2 FILE TOXCENTER
  - 220 FILE USPATFULL
  - 22 FILE USPAT2
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    - 38 FILE WPINDEX
  - 23 FILES HAVE ONE OR MORE ANSWERS, 56 FILES SEARCHED IN STNINDEX
- L1 QUE LACTOCOCCUS AND SKIN AND PROBIOTIC

=> logoff

ALL L# QUERIES AND ANSWER SETS ARE DELETED AT LOGOFF LOGOFF? (Y)/N/HOLD:y

COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 0.71 0.94

FULL ESTIMATED COST